This document is scheduled to be published in the Federal Register on 02/24/2022 and available online at federalregister.gov/d/2022-03901, and on govinfo.gov

DEPART

Drug Enforcement Administration

[Docket No. DEA-956]

Bulk Manufacturer of Controlled Substances Application: Johnson Matthey Pharmaceutical Materials Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Johnson Matthey Pharmaceutical Materials Inc. has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application on or before [INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on November 5, 2021, Johnson Matthey Pharmaceutical Materials Inc., 25 Patton Road, Devens, Massachusetts 01434, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s)**:**

Controlled Substance	Drug Code	Schedule

Amphetamine	1100	II
Methylphenidate	1724	II
Nabilone	7379	II
Hydrocodone	9193	II
Levorphanol	9220	II
Thebaine	9333	II
Alfentanil	9737	II
Remifentanil	9739	II
Sufentanil	9740	II

The company plans to support its other manufacturing facilities located in West Deptford, New Jersey and Conshohocken, Pennsylvania with manufacturing and analytical testing. In reference to drug code 9333 as bulk, the company plans to manufacture a Thebaine derivative for distribution to its customers. No other activities for these drug codes are authorized for this registration.

Matthew J. Strait,
Deputy Assistant Administrator.

[FR Doc. 2022-03901 Filed: 2/23/2022 8:45 am; Publication Date: 2/24/2022]